P.06

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REMARKS

It is noted that an error appears in this specification of a clerical/typographical nature, as more fully described in the amendment. The error occurred in good faith. Correction thereof does not involve such changes that would constitute new matter. Please enter the described amendments.

Respectfully submitted,

Dated: 23 Jan 2002

Bv

Reg. No. 44,854

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JAN 23 2002

TECHNOLOGY CENTER 2800



Application Serial No	09/945,393
Filing Date	August 30, 2001
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Title: Dielectric Material Forming Methods and Enhanced Dielectric Materials	

VERSION WITH MARKINGS TO SHOW CHANGES MADE ACCOMPANYING SUPPLEMENTAL PRELIMINARY AMENDMENT

In the Specification

The replacement specification paragraphs incorporate the following amendments.

<u>Underlines</u> indicate insertions and strikeouts indicate deletions.

Page 9, paragraph number [0025], and page 11-12, paragraph numbers [0030] and [0031] have been amended as follows:

Although the principles described herein are indicated as particularly applicable to TA₂O₅Ta₂O₅ dielectric materials, the invention may be further applicable to other dielectric materials containing tantalum and oxygen, as well as dielectric materials not containing tantalum and/or oxygen. Accordingly, in another aspect of the invention, a dielectric forming method includes chemisorbing a first dielectric material on a substrate and chemisorbing a second dielectric material on the first material, one of the first and second dielectric materials comprising oxygen and a Group IB to VIIIB element. An enhanced dielectric material can be formed containing the first and second dielectric materials. The enhanced dielectric can exhibit a dielectric constant greater that of the first dielectric material.

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method includes atomic layer depositing an oxide of Group IVB metal on a first dielectric material containing TA₂Θ₆Ta₂O₅ and forming a second dielectric material containing the chemisorbed oxide and the first dielectric material. As one example, the atomic layer depositing can include chemisorbing at least one Group IVB metal precursor on the first dielectric material followed by purging chemisorption byproducts and excess metal precursor from over the substrate. Exemplary precursors include tetrakis dimethyl amido titanium (TDMAT), zirconium t-butoxide, and other suitable materials as known to those skilled in the art. The metal precursors can be used alone or in combination. For example, titanium and zirconium could be deposited together. In processes where tantalum oxide is also formed by atomic layer depositing, tantalum ethoxy (Taeto) is one example of potentially several suitable precursors.

precursor on the chemisorbed Group IVB metal or tantalum and purging chemisorption byproducts and excess oxygen precursor from over the substrate. A chemisorption product of the Group IVB metal precursor and the oxygen precursor can comprise Group IVB metal oxide. A chemisorption product of the tantalum precursor and the oxygen precursor and the oxygen precursor can comprise a tantalum oxide, for example, $\mp A_2 O_5 Ta_2 O_5$. H₂O is one example of potentially several suitable oxygen precursors. However, a more preferable oxygen precursor will be of a type that does not oxidize silicon during ALD.

END OF DOCUMENT